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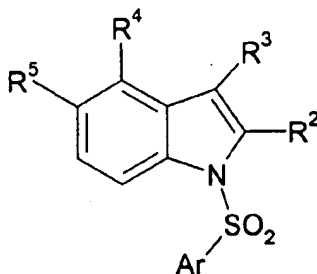
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(54) Title: 2-, 3-, 4-, OR 5-SUBSTITUTED-N1-(BENZENSULFONYL)INDOLES AND THEIR USE IN THERAPY

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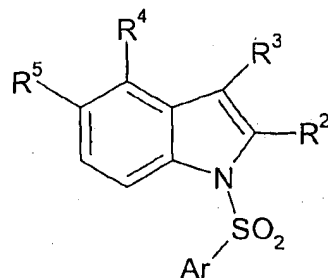


(I)

(57) Abstract: The invention provides 2, 3-, 4- or 5-substituted-N1-(ben-
zensulfonyl)indoles of the general formula (I) in which Ar, R²,
R³, R⁴ and R⁵ are as defined in the specification. The said compounds have
affinity for the 5-HT₂ receptor and are useful for the treatment and prophyl-
axis of disorders relating to the said receptor, such as obesity and CNS
disorders.

CLAIMS

1. A compound of formula (I):



(I)

5 wherein

Ar is

(1) phenyl,

(2) naphthyl,

- 10 (3) a 5- to 10-membered monocyclic or bicyclic heterocyclic ring having 1 to 4 heteroatoms selected from the group consisting of oxygen, sulfur, or nitrogen, or

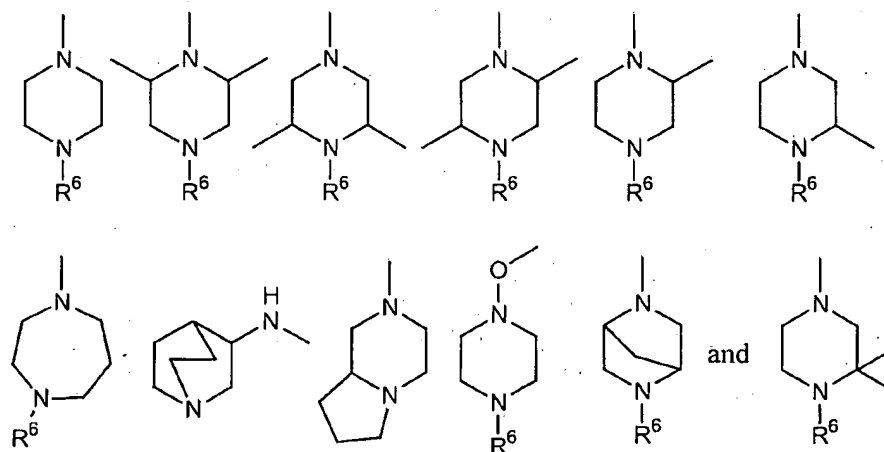
(4) -R⁹-phenyl;

- wherein each of phenyl, naphthyl, and heterocyclic ring is independently optionally substituted with halogen, C₁₋₆ alkyl, CF₃, hydroxyl, C₁₋₆ alkoxy, OCF₃, COCF₃, CN, NO₂, phenyloxy, phenyl, C₁₋₆ alkylsulfonyl, C₂₋₆ alkenyl, -NR⁷R⁸, C₁₋₆ alkylcarboxyl, formyl, -C₁₋₆ alkyl-NH-CO-phenyl, -C₁₋₆ alkyl-CO-NH-phenyl, -NH-CO-C₁₋₆ alkyl, -CO-NR⁷R⁸, or SR⁷; wherein each of R⁷ and R⁸ is independently H or C₁₋₆ alkyl; and R⁹ is C₁₋₆ alkyl or C₂₋₆ alkenyl, each of which being optionally substituted with phenyl or phenyloxy;

- 20 R² is H, phenyl, I, or C₁₋₆ alkyl;

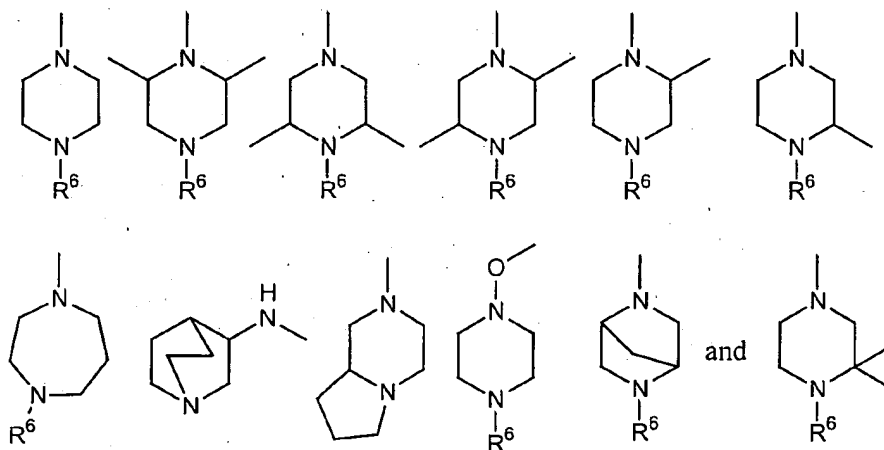
R³ is H or 3-(1-azabicyclo[2.2.2]oct-2-en)yl;

R⁴ is H or a heterocyclic ring selected from the group consisting of:



wherein R^6 is H, C_{1-6} alkyl, or benzyl; and

R^5 is H, hydroxy, C_{1-3} alkoxy, F, NO_2 , CF_3 , OCF_3 , or a heterocyclic ring selected from the group consisting of:



or a pharmaceutically acceptable salt, hydrate, or stereoisomer thereof,
with the proviso that when R^2 is alkyl, R^4 is not H.

2. A compound according to claim 1, wherein

Ar is

(1) phenyl,

(2) 1-naphthyl or 2-naphthyl,

(3) a 5- to 10-membered monocyclic or bicyclic heterocyclic ring having 1 to 4 heteroatoms selected from the group consisting of oxygen, sulfur, or nitrogen, or

(4) $-R^9$ -phenyl;

wherein each of phenyl, naphthyl, and heterocyclic ring is independently optionally substituted with F, Cl, Br, C_{1-6} alkyl, CF_3 , hydroxyl, C_{1-6} alkoxy, OCF_3 , phenyl, C_{2-6}

INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER

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B. FIELDS SEARCHED

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9636611 A1 (CHIROSCIENCE LIMITED), 21 November 1996 (21.11.96) --	1, 3, 10, 21-22
X	WO 9805315 A1 (TULARIK, INC.), 12 February 1998 (12.02.98) --	1, 3, 10, 21, 22
X	WO 9633171 A1 (ISTITUTO SUPERIORE DI SANITA'), 24 October 1996 (24.10.96) --	1, 3, 10, 21, 22

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

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Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, File CAPLUS, CAPLUS accession no.1993:539117, document no. 119:139117, Pfizer Inc., USA: "Heteroaryl amines as novel acetyl cholinesterase inhibitors"; & WO,9307140,A1,19930415 5577046 1752	1,3,10,21,22
X	STN International, File CAPLUS, CAPLUS accession no.1996:736044, document no. 126:29025, Bojinov, V. et al: "Synthesis of new N-arylsulfonylindoles and in vitro assay for fungicidal activity"; & Biotechnol. Biotechnol. Equip. (1996), (1), 27-31 --	1,3,10
X	STN International, File CAPLUS, CAPLUS accession no.1998:724128, document no. 130:21742, Nippon Soda Co., LTD.: "Pyridylindole compounds and agrochemical fungicides containing them", & JP,10298011,A2,19981110 --	1,3
X	STN International, File CAPLUS, CAPLUS accession no.1993:495527, document no. 119:95527, Mitsubishi Petrochemical Co., LTD.: "Preparation of carbamoyltriazole derivatives as herbicides"; & JP,04321671,A2,19921111 --	1,5,10
X	STN International, File CAPLUS, CAPLUS accession no.1993:147461, document no. 118:147461, Taisho Pharmaceutical Co., LTD.: "N-Phenylsulfonylindole derivatives"; & JP,04273857,A2,19920930 --	1,3,10,21,22

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C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>STN International, File CAPLUS, CAPLUS accession no.1999:550761, document no. 131:299130, Nyasse, Barthelemy et al: "2-Naphthalensulfonyl as a Tosyl Substitute for Protection of Amino Functions. Cyclic Voltammetry Studies on Model Sulfonamides and Their Preparative Cleavage by Reduction"; & J.Org.Chem. (1999), 64(19), 7135-7139</p> <p>--</p>	1,3,4,10
X	<p>STN International, File CAPLUS, CAPLUS accession no.1986:110010, document no. 104:110010, Ketcha, Daniel M. et al: "A convenient synthesis of 3-acylindoles via Friedel Crafts acylation of 1-(phenyl- sulfonyl)indole. A new route to pyridocarbazole- 5,11-quinones and ellipticine"; & J. Org. Chem. (1985), 50(26), 5451-7</p> <p>--</p>	1,3,10
X	<p>STN International, File CAPLUS, CAPLUS accession no.1979:450235, document no. 91:20235, Illi, Volker O.: "Phase transfer-catalyzed N-sulfonation of indole"; & Synthesis (1979), (2), 136</p> <p>--</p>	1,3,10
X	<p>STN International, File CAPLUS, CAPLUS accession no.1982:562734, document no. 97:162734. Obafemi, Craig A.: "Studies in the heterocyclic compounds. V. Some reactions of 5-chloro-2-thiophenesulfonyl derivatives", Phosphorus Sulfur (1982), 13(1),119-31</p> <p>--</p>	1,5,10
X	<p>STN International, File CAPLUS, CAPLUS accession no.1986:186352, document no. 104:186352, Obafemi, Craig A. et al: "Studies of heterocyclic compounds. 8. The synthesis and some reactions of 4- bromoimidazole-5-sulfonyl derivatives", J. Chem. Eng. Data (1986), 31(2), 257-9</p> <p>--</p>	1,5,10

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International application No.

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C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, File CAPLUS, CAPLUS accession no. 1987-156217, document no. 106:156217, Kasahara, Akira et al: "Palladium-catalyzed synthesis of 2- substituted indoles"; & Yamagata Daigaku Kiyo, Kogaku (1986), 19(1), 39-51 --	1,3,10
X	STN International, File CAPLUS, CAPLUS accession no. 1993:101757, document no. 118:101757, Fuji, Masahiro et al: "Preparation of alkyl-substituted indoles in the benzene portion. Part 6. Synthetic procedure for 4-,5-,6-, or 7-alkoxy- and hydroxyindole derivatives"; & Chem. Pharm. Bull. (1992), 40(9), 2344-52 --	1,3,10
X	STN International, File CAPLUS, CAPLUS accession no. 1995:751092, document no. 123:198572, Goulaouic-Dubois et al: "Protection of Amines by the Pyridine- 2-sulfonyl Group and Its Cleavage under Mild Conditions (SmI2 or Electrolysis); & J. Org. Chem. (1995), 60(18), 5969-72 --	1,5,10
X	STN International, File CAPLUS, CAPLUS accession no. 1998:265011, document no. 129:54315, Pathak, Vijai N. et al: "Fluorophenylsulfonylation of 4,5-dihydro- 3,5-diarylpyrazoles and 2-arylindoles via solid-liquid phase transfer catalysis"; & Indian J. Heterocycl. Chem. (1998), 7(3), 241-242 --	1,3

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International application No.

PCT/SE 01/02319

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, File CAPLUS, CAPLUS accession no. 1999:661842, document no. 132:35576, Ishikura, Minoru et al: "Investigation of the reaction of N-substituted indolylborates: palladium catalyzed cross-coupling reactions and intramolecular alkyl migration reactions"; & J. Heterocycl. Chem. (1999), 36(4), 873-879 --	1,3
X	STN International, File CAPLUS, CAPLUS accession no. 1998:710510, document no. 130:52304, Danieli, Bruno et al: "Application of the Pd-catalyzed heteroarylation of the synthesis of 5-(indol-2'-yl)pyridin-2-one and 5-(indol-2'-yl)pyran-2-one"; & Tetrahedron (1998), 54 (46), 14081-14088 --	1,3,10
X	STN International, File CAPLUS, CAPLUS accession no. 1999:235586, document no. 130:352114, Li, Zhaopeng et al: "Hetero- cyclic aromatic amide protecting groups for aryl and phthalocyaninesulfonic acids"; & Can. J. Chem. (1999), 77(1), 138-145 --	1,3,10
P,X	STN International, File CAPLUS, CAPLUS accession no. 2001:419497, document no. 135:226906, Merlic C. A. et al: "Benzannulation reactions of Fischer carbene complexes for the synthesis of indolocarbazoles"; & Tetrahedron (2001), 57(24), 5199-4020 --	1,3
P,X	STN International, File CAPLUS, CAPLUS accession no. 2001:466985, document no. 135:273099, Xiong, W.-N., et al: "Synthesis of Novel Analogues of Marine Indole Alkaloids: Mono(indolyl)-4-trifluoromethylpyridines and Bis (indolyl)-4-trifluoromethylpyridines as Potential Anticancer Agents"; & Biorganic & Medicinal Chemistry (2001), 9(7), 1773-1780 --	1,3,10